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## REISSUE PATENT APPLICATION TRANSMITTAL

Address to:		Attomey Docket No.	3254:1002-028	
Assistant Commissioner for Patents		First Named Inventor	Breslow	
Box Reissue		Original Patent Number Original Patent Issue Date	5,369,108	
Washington, DC 20231		(Month/Day/Year)	Nov. 29, 1994	
			Express Mail Label No.	EV 044389584 US
APPLICATION FOR REISSUE OF: (Check applicable box)  Utility Patent		Design Patent	Plant Patent	
APPLICATION ELEMENTS (37 CFR 1.173)		ACCOMPANYING APPLICATION PARTS		
	Fee Transmittal Form (PTO/ SB/ 56) (Submit an original, and a duplicate for fee processing)		10. X Statement of status and support for all changes to the claims. See 37 CFR 1.173 (c).	
2. Applicant clair	nt claims small entity status. See 37 CFR 1.27.		11. Original U.S. Patent for surrender	
	and Claims in double column copy o	f patent	Ribboned Origina	al Patent Grant
	Drawing(s) (proposed amendments, if appropriate)		Statement of Loss (PTO/SB/55)	
5. Reissue Oath/Declaration (original or copy) (2 docs – (37 C.F.R. § 1.175) (PTO/SB/51 or 52) 10 pgs total)		12. Foreign Priority Claim (35 U.S.C. 119) (if applicable)		
`	mey (included in Decl		13. X Information Disclos Statement (IDS)/PT	IX I F I
7. Original U.S. Patent currently assigned? X Yes No			14. English Translation of Reissue Oath/Declaration (if applicable)	
(If Yes, check applicable box(es))			(п аррпсаые)	
X Written Consent of all Assignees (PTO/SB/53) (2 docs)		15. X Preliminary Amendment		
X 37 C.F.R. § 3.73(b) Statement (2 docs) (PTO/SB/96)		16. X Return Receipt Postcard (MPEP 503) (Should be specifically itemized)		
8. CD-ROM or CD-R in duplicate, Computer Program (Appendix) or large table		17. Other:Checkfar\$.1994		
Nucleotide and/or Amino Acid Sequence Submission     (if applicable, all of the following are necessary)				
a. Computer Readable Form (CFR)				
b. Specification Sequence Listing on: i ☐ CD-ROM (2 copies) or CD-R (2 copies); <b>or</b> ii ☐ paper				
c. Statements verifying identity of above copies				
18. CORRESPONDENCE ADDRESS				
Customer Number or Bar Code Label 021005 or Correspondence address below (Insert Customer No. or Attach bar code label here)				
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NAME (PrintType) Susan M. Abelleira Ro		Registration No. (Attorney/Agent)	2,252	
Signature Susan M. Abelleura:				ovember 02,2001

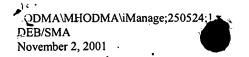
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Approved for use through 01/31/2004. OMB 0651-0033 U.S. Patent and Trademark Office; U.S. DEPARTMENT OF COMMERCE Under the Paperwork Reduction Act of 1995, no persons are required to respond to a collection of information unless it displays a valid OMB control number. Docket Number (Optional) 3254.1002-028 REISSUE APPLICATION FEE TRANSMITTAL FORM Claims as Filed - Part 1 Other than a Small Entity Small Entity Claims in (3) Number Filed in Patent Rate Fee Rate Fee Reissue Application Number Extra Total Claims 18. 162. (A) 17 (B) 9 29 (37 CFR 1.16(i)) or (C) 1 Independent claims 14 13 1092. x\$<u>84</u>= (37 CFR 1.16(i)) \$740. Basic Fee (37 CFR 1.16(h)) Total Filing Fee \$1994. \$ OR Claims as Amended - Part 2 (3) (2)(1) Small Entity Other than a Small Entity Highest Number Extra Claims Remaining Rate Fee Rate Fee Claims Previously After Amendment Paid For Present Total Claims = MINUS x \$. (37 CFR 1.16(j) Independent MINUS x \$ Claims (37 CFR 1.16(i)) \$ OR \$ **Total Additional Fee** \* If the entry in (D) is less than the entry in (C), Write "0" in column 3. \*\* If the "Highest Number of Total Claims Previously Paid For" is less than 20, Write "20" in this space. \*\*\* After any cancellation of claims. \*\*\*\* If "A" is greater than 20, use (B - A); if "A" is 20 or less, use (B - 20). \*\*\*\*\* "Highest Number of Independent Claims Previously Paid For" or Number of Independent Claims in Patent (C). Applicant claims small entity status. See 37 CFR 1.27. in the amount of . Please charge Deposit Account No. A duplicate copy of this sheet is enclosed. The Commissioner is hereby authorized to charge any additional fees under 37 CFR 1.16 or 1.17 which may be required, or credit any overpayment to Deposit Account No. \_\_\_\_\_08-0380 A duplicate copy of this sheet is enclosed. to cover the filing / additional fee is enclosed. X A check in the amount of \$ 1994 Payment by credit card. Form PTO-2038 is attached. WARNING: Information on this form may become public. Credit card information should not be included on this form. Provide credit card information and authorization on PTO-2038. Susan M. abelleura, 42,252
Signature of Applicant, Attorney or Agent of Record Movember 02, 2001 Susan M. Abelleira Typed or printed name



### IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicants:

Ronald Breslow, Paul A. Marks, Richard A. Rifkind, and Branko Jursic

Reissue Application of U.S. Patent No.: 5,369,108

Issue Date: November 29, 1994

For:

POTENT INDUCERS OF TERMINAL DIFFERENTIATION AND METHODS OF USE THEREOF

Date: 1/2/01

EXPRESS MAIL LABEL NO. EV 044389584 US

# STATUS OF CLAIMS AND SUPPORT FOR CLAIM CHANGES UNDER 37 C.F.R. §1.173(c)

Box REISSUE
Assistant Commissioner for Patents
Washington, D.C. 20231

Sir:

This paper is being filed in accordance with 37 C.F.R. §1.173(c) to provide the status of all patent claims, and of all added claims in view of the Preliminary Amendment being filed concurrently. In addition, an explanation of the support in the disclosure of the patent for the changes made to the claims as set forth in the Preliminary Amendment, is provided.





As of the date of the Preliminary Amendment being filed concurrently herewith, originally issued Claims 1-17 are pending and newly added Claims 18-29 are also pending.

### SUPPORT FOR CLAIM CHANGES

Claim 1 has been amended to recite that  $R_1$  and  $R_2$  of the structure set forth in the claim are the same and are a substituted or unsubstituted thiazoleamino group. Support for this amendment can be found, *inter alia*, in the originally issued claims and at Col. 7, lines 40-52 of the specification.

Claim 2 has been amended to be in independent form and to recite that  $R_2$  is a hydroxylamino group and is different from  $R_3$ -N- $R_4$ . In addition, the definition of n has been amended to recite that n is an integer from 5 to about 8. Support for these amendments can be found, *inter alia*, in the originally issued claims, at Col. 2, line 65-Col. 3, line 6, at Col. 6, line 59-Col. 7, line 7 and in Table 1, as Entries 2-5 of Column "CPD".

Claim 3 has been amended to further define the integer n as 6. Support for this amendment can be found in originally issued Claim 3 and throughout the specification.

Claims 5 and 7 have been amended to correct obvious typographical errors. Support for this amendment can be found, for example, in Table 1, Entries 51-53 and in the art.

Claims 11, 12 and 13 have been amended to depend from Claim 2 rather than Claim 3. In addition, Claim 11 has been amended to designate the pyridine as a gamma pyridine,  $\gamma$ -pyridine, rather than a delta pyridine,  $\delta$ -pyridine. This is an obvious error, as  $\delta$ -pyridine cannot be a substituent. Support for this amendment can be found in the structures at Col. 17, lines 55-62 and Col. 18, lines 25-33.

Claims 18-29 are newly added. These claims have been added to more specifically claim particular compounds set forth in the specification. Claim 18 is directed to the compositions of the structure set forth in the claim, wherein n is from 5 to about 8. Support for Claim 18 can be found, for example, in Table 1 as Entries 2-5. Claim 19 is directed to a pharmaceutical composition comprising a therapeutically effective amount of a compound having the structure



set forth in the claim wherein n is from 5 to about 8. Support for Claim 19 can be found, for example, in Table 1 as Entries 2-5 and in the specification at Col. 6, lines 28-31.

Newly added independent Claim 20 is directed to the compound set forth as Entry 3 in Table 1. As such, support for newly added Claim 20 can be found in Table 1 and also at Col. 26, lines 50-68. Newly added independent Claim 21 is directed to a pharmaceutical composition comprising a therapeutically effective amount of the compound set forth as Entry 3 in Table 1 and a pharmaceutically acceptable carrier. As such, support for newly added Claim 21 can be found in Table 1, at Col. 6, lines 28-31 and at Col. 26, lines 50-68 of the specification.

Newly added independent Claim 22 is directed a compound of the general formula set forth, for example, at Col. 6, line 60 and up of the specification, wherein R<sub>2</sub> is a hydroxylamino group, n is 6, R<sub>3</sub> is a γ-pyridine group and R<sub>4</sub> is hydrogen. Support for newly added Claim 22 can be found, *inter alia*, in originally issued Claim 11 and in the structures at Col. 17, lines 55-62 and Col. 18, lines 25-33. Newly added independent Claim 23 is directed to a pharmaceutical composition comprising a therapeutically effective amount of the compound described above with regard to Claim 22 and a pharmaceutically acceptable carrier. Support for newly added Claim 23 can be found, *inter alia*, in originally issued Claim 11 and in the structures at Col. 17, lines 55-62 and Col. 18, lines 25-33, and at Col. 6, lines 28-31 of the specification.

Newly added independent Claim 24 is directed to a compound of the general formula set forth, for example, at Col. 6, line 60 and up of the specification, wherein  $R_2$  is a hydroxylamino group, n is 6,  $R_3$  is  $\beta$ -pyridine group and  $R_4$  is hydrogen. Support for newly added Claim 24 can be found, *inter alia*, in originally issued Claim 12. Newly added independent Claim 25 is directed to a pharmaceutical composition comprising a therapeutically effective amount of the compound described above with regard to Claim 24, and a pharmaceutically acceptable carrier. Support for newly added Claim 25 can be found, *inter alia*, in originally issued Claim 12 and at Col. 6, lines 28-31 of the specification.

Newly added independent Claim 26 is directed to a compound of the general formula set forth, for example, at Col. 6, line 60 and up of the specification, wherein  $R_2$  is a hydroxylamino group, n is 6,  $R_3$  is  $\alpha$ -pyridine group and  $R_4$  is hydrogen. Support for newly added Claim 24 can be found, *inter alia*, in originally issued Claim 13. Newly added independent Claim 27 is directed to a pharmaceutical composition comprising a therapeutically effective amount of the



compound described above with regard to Claim 26 and a pharmaceutically acceptable carrier. Support for newly added Claim 27 can be found, *inter alia*, in originally issued Claim 13 and at Col. 6, lines 28-31 of the specification.

Newly added independent Claim 28 is directed to the compound set forth as Entry 19 in Table 1. As such, support for newly added Claim 28 can be found in Table 1. Newly added independent Claim 29 is directed to a pharmaceutical composition comprising a therapeutically effective amount of the compound set forth as Entry 19 in Table 1 and a pharmaceutically acceptable carrier. As such, support for newly added Claim 29 can be found in Table 1 and at Col. 6, lines 28-31 of the specification.

#### **CONCLUSION**

In view of the above, it is believed that the requirements of 37 C.F.R. § 1.173(c) are fulfilled. Further, it is believed that all pending claims (Claims 1-29) are in condition for allowance, and it is respectfully requested that the application be passed to issue. If the Examiner feels that a telephone conference would expedite prosecution of this case, the Examiner is invited to call the undersigned at (978) 341-0036.

Respectfully submitted,

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